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DISHWASHING PRODUCT

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Cross-Reference to Related Applications

15 This application is a division of allowed U.S. Patent Application Serial No. 09/994,564, filed on November 27, 2001, which is incorporated by reference herein.

Technical field

20 The present invention is in the field of dishwashing, in particular it relates to dishwashing methods including methods for washing dishware/tableware in an automatic dishwashing machine using dishwashing products in unit dose and especially pouch form. The methods of the invention are especially useful for the removal of cooked-, baked- and burnt-on soils from cookware and tableware.

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Background of the invention

Unitised doses of dishwashing detergents are found to be more attractive and convenient to some consumers because they avoid the need of the consumer to measure the product
30 thereby giving rise to a more precise dosing and avoiding wasteful overdosing or underdosing. For this reason automatic dishwashing detergent products in tablet form have become very popular. Detergent products in pouch form are also known in the art.

It is normally the objective of the detergent formulator chemist to optimise the amount of
35 actives delivered to the wash for a given unit cost. The amount of actives delivered to the

5 wash is, among other factors, determined by the shape, size and density of the unitised dose form.

One of the drawbacks of unitised dose form such as tablets is that they have a fixed shape. The shape of dishwashing machine dispensers, on the other hand, is different from
10 manufacturer to manufacturer. Tablets are designed to have a size and shape which fit all machine dispensers, this fact together with the mechanical properties of tablets usually constrains the amount of product composition which can be incorporated in the tablet. Similar considerations can also apply in the case of detergent products in pouch form.

15 Another drawback of detergent tablets is the fact that their manufacturing process requires the additional step of powder compaction. This slows down the dissolution rate of the ingredients forming the tablet, or requires the use of complex and expensive disintegrant systems, or makes it difficult to achieve differential dissolution of the detergent active ingredients.

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Another factor that can contribute to the inefficient delivery of actives to the wash, in the case of tablets, is the need for adding carrier materials, as for example porous materials able to bind active liquid materials, binders and disintegrants. In particular, the incorporation of liquid surfactants to powder form detergent compositions can raise
25 considerable processing difficulties and also the problem of poor dissolution through the formation of surfactant gel phases.

There is still the need for a unitised dose form which allows for optimum delivery of active components across different washing machine types and which provides improved
30 processing and dissolution characteristics.

Summary of the invention

35 According to a first aspect of the present invention, there is provided a method of washing dishware/tableware in an automatic dishwashing machine having a single or multi-

5 compartment product dispenser which is normally closed and sealed after charging the machine and prior to delivery of the dishwashing product into the wash liquor and wherein the dishwashing product comprises one or more dishwashing compositions in a unit dose form. The unit dose forms used herein are deformable and preferably have a shape and size such that they are compressibly contained within the product dispenser.

10 The dishwashing product has a deformability, as measured following the method described hereinbelow of greater than about 5%, preferably greater than about 8%, more preferably greater than about 10% and even more preferably greater than about 20%. The shape and size of the product are also such that it occupies at least about 60%, preferably at least about 70%, more preferably at least about 80%, especially more than about 85%

15 of the volume of the corresponding compartment of the product dispenser in its closed state. Provided that in the case of single compartment dispensers the dishwashing product can occupy at least about 40%, preferably at least about 50% of the volume of the product dispenser compartment in its closed state. The term "compressibly contained" as used herein means that the product is in a state of compression within the closed product

20 dispenser across at least one transverse section of the product. Preferably the product is in a state of compression across the smallest transverse section of the product in a direction generally perpendicular to the product dispenser closure means.

The deformability of the unit dose form may be measured using an Instron materials

25 tester (or similar) according to the following procedure. The unit dose form is placed on a flat surface such that it lies on a base of maximum footprint and a corresponding flat probe is brought down upon the upper surface of the unit dose form. The movement of the probe is continued until a sufficient reaction force is created to cause the unit dose form to fracture or burst. The deformability of the unit dose form may be defined as:

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$$\left\{ \frac{\text{displacement of probe after touching unit dose form up to burst point}}{\text{total unit dose form thickness}} \right\} \times 100$$

The volume of the unit-dose containing product dispenser compartment in its closed state lies in the range from about 15 to about 70, preferably from about 18 to 50 and more

35 preferably from about 20 to 30 ml. In the case of multi-compartment dispensers, the

5 individual compartments generally have a volume of from about 10 to about 35 ml, preferably from about 15 to about 30 ml. The total volume of the product dispenser (both multi and single compartments) on the other hand is generally from about 20 to about 70 ml, preferably from about 30 to about 50 ml.

10 The deformability herein is measured when the unit dose form is resting on its maximum footprint. The deformability measured when the pouch is placed in this position is sometimes referred to herein as "vertical deformability". The deformability measured when the unit dose form is rotated into a perpendicular plane is referred herein as "horizontal deformability". In preferred embodiments the unit doses have differing
15 vertical and horizontal deformability (so-called anisotropic deformability), the vertical and minimum horizontal deformability in the perpendicular plane preferably differing from each other by at least about 30% preferably at least about 40%. It is also preferred that the minimum horizontal deformability is greater than the vertical deformability. Anisotropic deformability is preferred herein from the dispenser fit, packaging and the
20 feel and handling viewpoints.

The term "unit dose" herein refer to a dose of detergent product incorporating one or more dishwashing compositions and sufficient for a single wash cycle. Suitable unit dose forms include capsules, sachets and pouches which can have single or multiple
25 compartments. Suitable unit dose forms for use herein include water-soluble, water-dispersible and water-permeable capsules, sachets and pouches. Preferred for use herein are water soluble pouches, based on partially hydrolysed polyvinyl alcohol as pouch material. Dishwashing compositions incorporated therein can be in liquid, gel, paste or pouch form, but preferably composition in liquid gel or paste form are substantially
30 anhydrous for reasons of pouch stability.

In a preferred aspect of the invention, dishwashing product comprises a dose sufficient for a single wash cycle of an anhydrous dishwashing composition. The term anhydrous as used herein is intended to include compositions containing less than about 10% of water
35 by weight of the composition, preferably less than about 5% of water and more preferably less than about 1%. The water can be present in the form of hydrated compounds, i.e.

5 bound water or in the form of moisture. It is preferred that the composition contains less than about 1%, preferably less than about 0.1% free moisture. Free moisture can be measured by extracting 2 g of the product into 50 ml of dry methanol at room temperature for 20 minutes and then analysis a 1 ml aliquot of the methanol by Karl Fischer titration.

10 In preferred embodiments the dishwashing composition comprises an organic solvent system. The organic solvent system can simply act as a liquid carrier, but in preferred compositions, the solvent can aid removal of cooked-, baked- or burnt-on soil and thus has detergent functionality in its own right. The organic solvent system (comprising a single solvent compound or a mixture of solvent compounds) preferably has a volatile
15 organic content above 1 mm Hg and more preferably above 0.1 mm Hg of less than about 50%, preferably less than about 20% and more preferably less than about 10% by weight of the solvent system. Herein volatile organic content of the solvent system is defined as the content of organic components in the solvent system having a vapor pressure higher than the prescribed limit at 25°C and atmospheric pressure.

20

The organic solvent system for use herein is preferably selected from organoamine solvents, inclusive of alkanolamines, alkylamines, alkyleneamines and mixtures thereof; alcoholic solvents inclusive of aromatic, aliphatic (preferably C₄-C₁₀) and cycloaliphatic alcohols and mixtures thereof; glycols and glycol derivatives inclusive of C₂-C₃
25 (poly)alkylene glycols, glycol ethers, glycol esters and mixtures thereof; and mixtures selected from organoamine solvents, alcoholic solvents, glycols and glycol derivatives. In one preferred embodiment the organic solvent comprises organoamine (especially alkanolamine) solvent and glycol ether solvent, preferably in a weight ratio of from about 3:1 to about 1:3, and wherein the glycol ether solvent is selected from ethylene glycol
30 monobutyl ether, diethylene glycol monobutyl ether, ethylene glycol monomethyl ether, ethylene glycol monoethyl ether, diethylene glycol monomethyl ether, diethylene glycol monoethyl ether, propylene glycol monobutyl ether, and mixtures thereof. Preferably, the glycol ether is a mixture of diethylene glycol monobutyl ether and propylene glycol butyl ether, especially in a weight ratio of from about 1:2 to about 2:1.

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5 In another embodiment of the invention, the anhydrous dishwashing composition is in the form of a particulate bleach suspension in a non-aqueous liquid carrier. Preferred liquid carriers comprises at least about 50%, preferably at least about 60% and more preferable at least 90% of a solvent or solvent mixture having:

- 10 i) a fractional dispersion Hansen solubility parameter greater than about 40%, preferably greater than about 60% and more preferably greater than about 80%; and
- ii) a fractional polar Hansen solubility parameter less than about 60%, preferably less than about 40% and more preferably less than about 20%.

Fractional dispersion Hansen solubility parameter of a solvent is defined as the ratio (multiplied by 100) of the dispersion Hansen solubility parameter to the sum of the
15 dispersion, polar and hydrogen bonding Hansen solubility parameters. Fractional polar Hansen solubility parameter of a solvent is accordingly defined.

Solvents having the fractional Hansen solubility parameters described hereinabove are particularly valuable for purposes of bleach stability. These solvents have very low water
20 absorption, this is particularly important in cases wherein the bleach is contained in pouches, because apart from the problem of loss of bleach, bleach decomposition gives rise to oxygen gas which can cause bloating of the pouch material and give the pouches a fluffy appearance (not very attractive to the consumers). Particulate bleaches suitable for use herein include inorganic peroxides inclusive of perborates and percarbonates, organic
25 peracids inclusive of preformed monoperoxy carboxylic acids, such as phthaloyl amido peroxy hexanoic acid and di-acyl peroxides. Preferred peroxides for use herein are percarbonate and perborate bleach.

One problem in formulating particulate bleach into liquid compositions is to keep the
30 bleach physically stable and homogeneously distributed in the liquid composition. Bleach suspension can be achieved by matching the density of the liquid carrier and the particulate bleach. To this end, the density difference between the particulate bleach and the non-aqueous liquid carrier is preferably less than about 500 Kg /m³, more preferably less than about 300 Kg /m³. High viscosity and small particle size will also contribute to
35 the formation of a stable suspension. In one embodiment of the present invention, the particulate bleach has an average particle size from about 10 µm to about 500 µm,

5 preferably from about 30 μm to about 250 μm , as measured using a Malvern particle size
analyser based on laser diffraction. The suitable viscosities for the suspensions of the
invention are from about 1,000 Kg/m s^{-1} to about 100,000 Kg/m s^{-1} , preferably from
about 5,000 Kg/m s^{-1} to about 50,000 Kg/m s^{-1} at shear rate of 1s^{-1} ; and from about 500
10 Kg/m s^{-1} to about 50,000 Kg/m s^{-1} , preferably from about 800 Kg/m s^{-1} to about 30,000
 Kg/m s^{-1} at shear rate of 150s^{-1} as measured using a Contraves Rheometer with 40 mm
diameter parallel plate at 25°C .

In preferred embodiments the dishwashing composition included in the unit dose form of
the invention comprises a detergent enzyme. In another embodiment the dishwashing
15 composition comprises an alkalinity source.

According to another aspect of the invention there is provided a unit dose dishwashing
detergent composition in the form of a paste having a density greater than about 1100
20 Kg/m^3 , preferably greater than about 1300 Kg/m^3 .

Multi-compartment pouches suitable for use herein can include compartments with
different solubilities controlled by for example pH, temperature or any other means. High
temperature soluble pouches allow the handling of the pouches at ambient temperature
with wet hands.

25 Unitised doses having multi-compartment can comprise at least one compartment
containing a powder composition. This powder composition comprises traditional solid
materials used in dishwashing detergent, such as builders, alkalinity sources, bleaches,
etc. Especially useful are multi-compartment unit dose forms comprising different
30 compartments for solid and for liquid compositions. The liquid compositions comprise
traditional liquid materials used in dishwashing detergents, such as non-ionic surfactants
or the organic solvents described hereinabove. Especially useful liquids for use in the
case of multi-compartment unit dose forms comprising a powder compartment and a
liquid compartment are liquids with hygroscopic and hydrophilic properties because they
35 are capable to act as a moisture sink and reduce moisture pick-up by the powder
compartment.

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Detailed description of the invention

The present invention envisages the use of dishwashing detergent composition in unit dose form, which have a high degree of deformability. This allow optimal use of the dishwashing machine dispenser, without losing the convenience of unit dose form. The invention also envisages the use of single and multi-compartment unit dose forms. Single compartment unit dose executions are especially useful in the case of paste/paste-like compositions. Multi-compartment unit dose form executions include unit dose forms comprising anhydrous liquids, especially useful compositions are those containing an organic solvent capable of remove baked-, cook- or burnt-on soils. The invention also envisages the use of anhydrous suspensions containing particulate bleach. Other forms of multi-compartment executions include a powder containing compartment in combination with a liquid containing compartment.

Unitised dose forms specially useful for use herein are pouches. The pouch herein is typically a closed structure which comprises one or more compartments, made of materials described herein. Subject to the constraints of deformability and dispenser fit, the pouch can be of any form, shape and material which is suitable to hold the composition, e.g. without allowing the release of the composition from the pouch prior to contact of the pouch to water. The exact execution will depend on, for example, the type and amount of the composition in the pouch, the number of compartments in the pouch, the characteristics required from the pouch to hold, protect and deliver or release the composition and/or components thereof.

The composition, or components thereof, are contained in the internal volume space of the pouch, and are typically separated from the outside environment by a barrier of water-soluble material. Typically, different components of the composition contained in different compartments of the pouch are separated from one another by a barrier of water-soluble material.

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5 In the case of multi-compartment pouches, the compartments may be of a different colour from each other, for example a first compartment may be green or blue, and a second compartment may be white or yellow. One compartment of the pouch may be opaque or semi-opaque, and a second compartment of the pouch may be translucent, transparent, or semi-transparent. The compartments of the pouch may be the same size, having the same
10 internal volume, or may be different sizes having different internal volumes.

For reasons of deformability and dispenser fit under compression forces, pouches or pouch compartments containing a component which is liquid will usually contain an air bubble having a volume of up to about 50%, preferably up to about 40%, more preferably
15 up to about 30%, more preferably up to about 20%, more preferably up to about 10% of the volume space of said compartment.

The pouch is preferably made of a pouch material which is soluble or dispersible in water, and has a water-solubility of at least 50%, preferably at least 75% or even at least 95%, as
20 measured by the method set out here after using a glass-filter with a maximum pore size of 20 microns.

50 grams \pm 0.1 gram of pouch material is added in a pre-weighed 400 ml beaker and 245ml \pm 1ml of distilled water is added. This is stirred vigorously on a magnetic stirrer
25 set at 600 rpm, for 30 minutes. Then, the mixture is filtered through a folded qualitative sintered-glass filter with a pore size as defined above (max. 20 micron). The water is dried off from the collected filtrate by any conventional method, and the weight of the remaining material is determined (which is the dissolved or dispersed fraction). Then, the % solubility or dispersability can be calculated.

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Preferred pouch materials are polymeric materials, preferably polymers which are formed into a film or sheet. The pouch material can, for example, be obtained by casting, blow-moulding, extrusion or blow extrusion of the polymeric material, as known in the art.

35 Preferred polymers, copolymers or derivatives thereof suitable for use as pouch material are selected from polyvinyl alcohols, polyvinyl pyrrolidone, polyalkylene oxides,

5 acrylamide, acrylic acid, cellulose, cellulose ethers, cellulose esters, cellulose amides, polyvinyl acetates, polycarboxylic acids and salts, polyaminoacids or peptides, polyamides, polyacrylamide, copolymers of maleic/acrylic acids, polysaccharides including starch and gelatine, natural gums such as xanthum and carragum. More preferred polymers are selected from polyacrylates and water-soluble acrylate
10 copolymers, methylcellulose, carboxymethylcellulose sodium, dextrin, ethylcellulose, hydroxyethyl cellulose, hydroxypropyl methylcellulose, maltodextrin, polymethacrylates, and most preferably selected from polyvinyl alcohols, polyvinyl alcohol copolymers and hydroxypropyl methyl cellulose (HPMC), and combinations thereof. Preferably, the level of polymer in the pouch material, for example a PVA polymer, is at least 60%.

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The polymer can have any weight average molecular weight, preferably from about 1000 to 1,000,000, more preferably from about 10,000 to 300,000 yet more preferably from about 20,000 to 150,000.

20 Mixtures of polymers can also be used as the pouch material. This can be beneficial to control the mechanical and/or dissolution properties of the compartments or pouch, depending on the application thereof and the required needs. Suitable mixtures include for example mixtures wherein one polymer has a higher water-solubility than another polymer, and/or one polymer has a higher mechanical strength than another polymer.
25 Also suitable are mixtures of polymers having different weight average molecular weights, for example a mixture of PVA or a copolymer thereof of a weight average molecular weight of about 10,000- 40,000, preferably around 20,000, and of PVA or copolymer thereof, with a weight average molecular weight of about 100,000 to 300,000, preferably around 150,000.

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Also suitable herein are polymer blend compositions, for example comprising hydrolytically degradable and water-soluble polymer blends such as polylactide and polyvinyl alcohol, obtained by mixing polylactide and polyvinyl alcohol, typically comprising about 1-35% by weight polylactide and about 65% to 99% by weight
35 polyvinyl alcohol.

5 Preferred for use herein are polymers which are from about 60% to about 98% hydrolysed, preferably about 80% to about 90% hydrolysed, to improve the dissolution characteristics of the material.

Most preferred pouch materials are PVA films known under the trade reference Monosol
10 M8630, as sold by Chris-Craft Industrial Products of Gary, Indiana, US, and PVA films of corresponding solubility and deformability characteristics. Other films suitable for use herein include films known under the trade reference PT film or the K-series of films supplied by Aicello, or VF-HP film supplied by Kuraray.

15 The pouch material herein can also comprise one or more additive ingredients. For example, it can be beneficial to add plasticisers, for example glycerol, ethylene glycol, diethyleneglycol, propylene glycol, sorbitol and mixtures thereof. Other additives include functional detergent additives to be delivered to the wash water, for example organic polymeric dispersants, etc.

20

The pouch can be prepared according to methods known in the art. The pouch is typically prepared by first cutting an appropriately sized piece of pouch material, preferably the pouch material. The pouch material is then folded to form the necessary number and size of compartments and the edges are sealed using any suitable technology,
25 for example heat sealing, wet sealing or pressure sealing. Preferably, a sealing source is brought into contact with the pouch material, heat or pressure is applied and the pouch material is sealed.

The pouch material is typically introduced to a mould and a vacuum applied so that the
30 pouch material is flush with the inner surface of the mould, thus forming a vacuum formed indent or niche in said pouch material. This is referred to as vacuum-forming.

Another suitable method is thermo-forming. Thermo-forming typically involves the step of forming an open pouch in a mould under application of heat, which allows the pouch
35 material to take on the shape of the mould.

5 Typically more than one piece of pouch material is used for making multi-compartment
pouches. For example, a first piece of pouch material can be vacuum pulled into the
mould so that said pouch material is flush with the inner walls of the mould. A second
piece of pouch material can then be positioned such that it at least partially overlaps, and
preferably completely overlaps, with the first piece of pouch material. The first piece of
10 pouch material and second piece of pouch material are sealed together. The first piece of
pouch material and second piece of pouch material can be made of the same type of
material or can be different types of material.

In a preferred process, a piece of pouch material is folded at least twice, or at least three
15 pieces of pouch material are used, or at least two pieces of pouch material are used
wherein at least one piece of pouch material is folded at least once. The third piece of
pouch material, or a folded piece of pouch material, creates a barrier layer that, when the
sachet is sealed, divides the internal volume of said sachet into at least two or more
compartments.

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The pouch can also be prepared by fitting a first piece of the pouch material into a mould,
for example the first piece of film may be vacuum pulled into the mould so that said film
is flush with the inner walls of the mould. A composition, or component thereof, is
typically poured into the mould. A pre-sealed compartment made of pouch material, is
25 then typically placed over the mould containing the composition, or component thereof.
The pre-sealed compartment preferably contains a composition, or component thereof.
The pre-sealed compartment and said first piece of pouch material may be sealed together
to form the pouch.

30 The detergent and cleaning compositions herein can comprise traditional detergency
components and can also comprise organic solvents having a cleaning function and
organic solvents having a carrier or diluent function or some other specialised function.
The compositions will generally be built and comprise one or more detergent active
components which may be selected from colorants, bleaching agents, surfactants,
35 alkalinity sources, enzymes, thickeners (in the case of liquid, paste, cream or gel
compositions), anti-corrosion agents (e.g. sodium silicate) and disrupting and binding

5 agents (in the case of powder, granules or tablets). Highly preferred detergent components include a builder compound, an alkalinity source, a surfactant, an enzyme and a bleaching agent.

10 Unless otherwise specified, the components described hereinbelow can be incorporated either in the organic solvent compositions and/or the detergent or cleaning compositions.

The organic solvents should be selected so as to be compatible with the tableware/cookware as well as with the different parts of an automatic dishwashing machine. Furthermore, the solvent system should be effective and safe to use having a
15 volatile organic content above 1 mm Hg (and preferably above 0.1 mm Hg) of less than about 50%, preferably less than about 30%, more preferably less than about 10% by weight of the solvent system. Also they should have very mild pleasant odours. The individual organic solvents used herein generally have a boiling point above about 150°C, flash point above about 100°C and vapor pressure below about 1 mm Hg, preferably
20 below 0.1 mm Hg at 25°C and atmospheric pressure.

Solvents that can be used herein include: i) alcohols, such as benzyl alcohol, 1,4-cyclohexanedimethanol, 2-ethyl-1-hexanol, furfuryl alcohol, 1,2-hexanediol and other similar materials; ii) amines, such as alkanolamines (e.g. primary alkanolamines:
25 monoethanolamine, monoisopropanolamine, diethylethanolamine, ethyl diethanolamine; secondary alkanolamines: diethanolamine, diisopropanolamine, 2-(methylamino)ethanol; ternary alkanolamines: triethanolamine, triisopropanolamine); alkylamines (e.g. primary alkylamines: monomethylamine, monoethylamine, monopropylamine, monobutylamine, monopentylamine, cyclohexylamine), secondary alkylamines: (dimethylamine), alkylene
30 amines (primary alkylene amines: ethylenediamine, propylenediamine) and other similar materials; iii) esters, such as ethyl lactate, methyl ester, ethyl acetoacetate, ethylene glycol monobutyl ether acetate, diethylene glycol monoethyl ether acetate, diethylene glycol monobutyl ether acetate and other similar materials; iv) glycol ethers, such as ethylene glycol monobutyl ether, diethylene glycol monobutyl ether, ethylene glycol monomethyl
35 ether, ethylene glycol monoethyl ether, diethylene glycol monomethyl ether, diethylene glycol monoethyl ether, propylene glycol butyl ether and other similar materials; v)

5 glycols, such as propylene glycol, diethylene glycol, hexylene glycol (2-methyl-2, 4 pentanediol), triethylene glycol, composition and dipropylene glycol and other similar materials; and mixtures thereof.

10 Surfactant

In the methods of the present invention for use in automatic dishwashing the detergent surfactant is preferably low foaming by itself or in combination with other components (i.e. suds suppressers). Surfactants suitable herein include anionic surfactants such as alkyl sulfates, alkyl ether sulfates, alkyl benzene sulfonates, alkyl glyceryl sulfonates, 15 alkyl and alkenyl sulphonates, alkyl ethoxy carboxylates, N-acyl sarcosinates, N-acyl taurates and alkyl succinates and sulfosuccinates, wherein the alkyl, alkenyl or acyl moiety is C₅-C₂₀ , preferably C₁₀-C₁₈ linear or branched; cationic surfactants such as chlorine esters (US-A-4228042, US-A-4239660 and US-A-4260529) and mono C₆-C₁₆ N-alkyl or alkenyl ammonium surfactants wherein the remaining N positions are 20 substituted by methyl, hydroxyethyl or hydroxypropyl groups; low and high cloud point nonionic surfactants and mixtures thereof including nonionic alkoxyated surfactants (especially ethoxylates derived from C₆-C₁₈ primary alcohols), ethoxylated-propoxylated alcohols (e.g., Olin Corporation's Poly-Tergent® SLF18), epoxy-capped poly(oxyalkylated) alcohols (e.g., Olin Corporation's Poly-Tergent® SLF18B - see WO-A-94/22800), 25 ether-capped poly(oxyalkylated) alcohol surfactants, and block polyoxyethylene-polyoxypropylene polymeric compounds such as PLURONIC®, REVERSED PLURONIC®, and TETRONIC® by the BASF-Wyandotte Corp., Wyandotte, Michigan; amphoteric surfactants such as the C₁₂-C₂₀ alkyl amine oxides (preferred amine oxides for use herein include lauryldimethyl amine oxide and hexadecyl 30 dimethyl amine oxide), and alkyl amphocarboxylic surfactants such as Miranol™ C2M; and zwitterionic surfactants such as the betaines and sultaines; and mixtures thereof. Surfactants suitable herein are disclosed, for example, in US-A-3,929,678 , US-A-4,259,217, EP-A-0414 549, WO-A-93/08876 and WO-A-93/08874. Surfactants are typically present at a level of from about 0.2% to about 30% by weight, more preferably 35 from about 0.5% to about 10% by weight, most preferably from about 1% to about 5% by

5 weight of composition. Preferred surfactant for use herein are low foaming and include low cloud point nonionic surfactants and mixtures of higher foaming surfactants with low cloud point nonionic surfactants which act as suds suppresser therefor.

10 Builder

Builders suitable for use in detergent and cleaning compositions herein include water-soluble builders such as citrates, carbonates and polyphosphates e.g. sodium tripolyphosphate and sodium tripolyphosphate hexahydrate, potassium tripolyphosphate and mixed sodium and potassium tripolyphosphate salts; and partially water-soluble or
15 insoluble builders such as crystalline layered silicates (EP-A-0164514 and EP-A-0293640) and aluminosilicates inclusive of Zeolites A, B, P, X, HS and MAP. The builder is typically present at a level of from about 1% to about 80% by weight, preferably from about 10% to about 70% by weight, most preferably from about 20% to about 60% by weight of composition.

20

Amorphous sodium silicates having an $\text{SiO}_2\text{:Na}_2\text{O}$ ratio of from 1.8 to 3.0, preferably from 1.8 to 2.4, most preferably 2.0 can also be used herein although highly preferred from the viewpoint of long term storage stability are compositions containing less than about 22%, preferably less than about 15% total (amorphous and crystalline) silicate.

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Enzyme

Enzymes suitable herein include bacterial and fungal cellulases such as Carezyme and Celluzyme (Novo Nordisk A/S); peroxidases; lipases such as Amano-P (Amano
30 Pharmaceutical Co.), M1 Lipase^R and Lipomax^R (Gist-Brocades) and Lipolase^R and Lipolase Ultra^R (Novo); cutinases; proteases such as Esperase^R, Alcalase^R, Durazym^R and Savinase^R (Novo) and Maxatase^R, Maxacal^R, Properase^R and Maxapem^R (Gist-Brocades); and α and β amylases such as Purafect Ox Am^R (Genencor) and Termamyl^R, Ban^R, Fungamyl^R, Duramyl^R, and Natalase^R (Novo); and mixtures thereof. Enzymes are
35 preferably added herein as prills, granulates, or cogranulates at levels typically in the range from about 0.0001% to about 2% pure enzyme by weight of composition.

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Bleaching agent

Bleaching agents suitable herein include chlorine and oxygen bleaches, especially inorganic perhydrate salts such as sodium perborate mono- and tetrahydrates and sodium percarbonate optionally coated to provide controlled rate of release (see, for example, GB-A-1466799 on sulfate/carbonate coatings), preformed organic peroxyacids and mixtures thereof with organic peroxyacid bleach precursors and/or transition metal-containing bleach catalysts (especially manganese or cobalt). Inorganic perhydrate salts are typically incorporated at levels in the range from about 1% to about 40% by weight, preferably from about 2% to about 30% by weight and more preferably from about 5% to about 25% by weight of composition. Peroxyacid bleach precursors preferred for use herein include precursors of perbenzoic acid and substituted perbenzoic acid; cationic peroxyacid precursors; peracetic acid precursors such as TAED, sodium acetoxymethylbenzene sulfonate and pentaacetylglucose; pernonanoic acid precursors such as sodium 3,5,5-trimethylhexanoyloxybenzene sulfonate (iso-NOBS) and sodium nonanoyloxybenzene sulfonate (NOBS); amide substituted alkyl peroxyacid precursors (EP-A-0170386); and benzoxazin peroxyacid precursors (EP-A-0332294 and EP-A-0482807). Bleach precursors are typically incorporated at levels in the range from about 0.5% to about 25%, preferably from about 1% to about 10% by weight of composition while the preformed organic peroxyacids themselves are typically incorporated at levels in the range from 0.5% to 25% by weight, more preferably from 1% to 10% by weight of composition. Bleach catalysts preferred for use herein include the manganese triazacyclononane and related complexes (US-A-4246612, US-A-5227084); Co, Cu, Mn and Fe bispyridylamine and related complexes (US-A-5114611); and pentamine acetate cobalt(III) and related complexes (US-A-4810410).

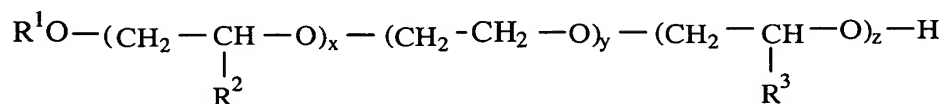
Low cloud point non-ionic surfactants and suds suppressers

The suds suppressers suitable for use herein include nonionic surfactants having a low cloud point. "Cloud point", as used herein, is a well known property of nonionic surfactants which is the result of the surfactant becoming less soluble with increasing

5 temperature, the temperature at which the appearance of a second phase is observable is referred to as the "cloud point" (See Kirk Othmer, pp. 360-362). As used herein, a "low cloud point" nonionic surfactant is defined as a nonionic surfactant system ingredient having a cloud point of less than 30° C., preferably less than about 20° C., and even more preferably less than about 10° C., and most preferably less than about 7.5° C. Typical
 10 low cloud point nonionic surfactants include nonionic alkoxyated surfactants, especially ethoxylates derived from primary alcohol, and polyoxypropylene/polyoxyethylene/polyoxypropylene (PO/EO/PO) reverse block polymers. Also, such low cloud point nonionic surfactants include, for example, ethoxylated-propoxylated alcohol (e.g., Olin Corporation's Poly-Tergent® SLF18) and
 15 epoxy-capped poly(oxyalkylated) alcohols (e.g., Olin Corporation's Poly-Tergent® SLF18B series of nonionics, as described, for example, in US-A-5,576,281).

Preferred low cloud point surfactants are the ether-capped poly(oxyalkylated) suds suppresser having the formula:

20



wherein R¹ is a linear, alkyl hydrocarbon having an average of from about 7 to about 12 carbon atoms, R² is a linear, alkyl hydrocarbon of about 1 to about 4 carbon atoms, R³ is a
 25 linear, alkyl hydrocarbon of about 1 to about 4 carbon atoms, x is an integer of about 1 to about 6, y is an integer of about 4 to about 15, and z is an integer of about 4 to about 25.

Other low cloud point nonionic surfactants are the ether-capped poly(oxyalkylated) having the formula:

30



wherein, R_I is selected from the group consisting of linear or branched, saturated or unsaturated, substituted or unsubstituted, aliphatic or aromatic hydrocarbon radicals having from about 7 to about 12 carbon atoms; R_{II} may be the same or different, and is

- 5 independently selected from the group consisting of branched or linear C₂ to C₇ alkylene in any given molecule; n is a number from 1 to about 30; and R_{III} is selected from the group consisting of:
- (i) a 4 to 8 membered substituted, or unsubstituted heterocyclic ring containing from 1 to 3 hetero atoms; and
 - 10 (ii) linear or branched, saturated or unsaturated, substituted or unsubstituted, cyclic or acyclic, aliphatic or aromatic hydrocarbon radicals having from about 1 to about 30 carbon atoms;
 - (b) provided that when R² is (ii) then either: (A) at least one of R¹ is other than C₂ to C₃ alkylene; or (B) R² has from 6 to 30 carbon atoms, and with the further
 - 15 proviso that when R² has from 8 to 18 carbon atoms, R is other than C₁ to C₅ alkyl.

Other suitable components herein include organic polymers having dispersant, anti-redeposition, soil release or other detergency properties in levels of from about 0.1% to about 30%, preferably from about 0.5% to about 15%, most preferably from about 1% to about 10% by weight of composition. Preferred anti-redeposition polymers herein include acrylic acid containing polymers such as Sokalan PA30, PA20, PA15, PA10 and Sokalan CP10 (BASF GmbH), Acusol 45N, 480N, 460N (Rohm and Haas), acrylic acid/maleic acid copolymers such as Sokalan CP5 and acrylic/methacrylic

25 copolymers. Preferred soil release polymers herein include alkyl and hydroxyalkyl celluloses (US-A-4,000,093), polyoxyethylenes, polyoxypropylenes and copolymers thereof, and nonionic and anionic polymers based on terephthalate esters of ethylene glycol, propylene glycol and mixtures thereof.

30 Heavy metal sequestrants and crystal growth inhibitors are suitable for use herein in levels generally from about 0.005% to about 20%, preferably from about 0.1% to about 10%, more preferably from about 0.25% to about 7.5% and most preferably from about 0.5% to about 5% by weight of composition, for example diethylenetriamine penta(methylene phosphonate), ethylenediamine tetra(methylene phosphonate)

35 hexamethylenediamine tetra(methylene phosphonate), ethylene diphosphonate, hydroxy-

5 ethylene-1,1-diphosphonate, nitrilotriacetate, ethylenediaminetetracetate,
ethylenediamine-N,N'-disuccinate in their salt and free acid forms.

The compositions herein can contain a corrosion inhibitor such as organic silver coating
agents in levels of from about 0.05% to about 10%, preferably from about 0.1% to about
10 5% by weight of composition (especially paraffins such as Winog 70 sold by Wintershall,
Salzbergen, Germany), nitrogen-containing corrosion inhibitor compounds (for example
benzotriazole and benzimidazole - see GB-A-1137741) and Mn(II) compounds,
particularly Mn(II) salts of organic ligands in levels of from about 0.005% to about 5%,
preferably from about 0.01% to about 1%, more preferably from about 0.02% to about
15 0.4% by weight of the composition.

Other suitable components herein include colorants, water-soluble bismuth compounds
such as bismuth acetate and bismuth citrate at levels of from about 0.01% to about 5%,
enzyme stabilizers such as calcium ion, boric acid, propylene glycol and chlorine bleach
20 scavengers at levels of from about 0.01% to about 6%, lime soap dispersants (see WO-A-
93/08877), suds suppressors (see WO-93/08876 and EP-A-0705324), polymeric dye
transfer inhibiting agents, optical brighteners, perfumes, fillers and clay.

Liquid detergent compositions can contain low quantities of low molecular weight
25 primary or secondary alcohols such as methanol, ethanol, propanol and isopropanol can
be used in the liquid detergent of the present invention. Other suitable carrier solvents
used in low quantities includes glycerol, propylene glycol, ethylene glycol, 1,2-
propanediol, sorbitol and mixtures thereof.

30 Examples:

Abbreviations used in Examples

In the examples, the abbreviated component identifications have the following meanings:

Carbonate : Anhydrous sodium carbonate

STPP (anhydrous) : Sodium tripolyphosphate anhydrous

STPP (hydrated)	:	Sodium tripolyphosphate hydrated to approximately 8%
Silicate	:	Amorphous Sodium Silicate ($\text{SiO}_2:\text{Na}_2\text{O}$ = from 2:1 to 4:1)
HEDP	:	Ethane 1-hydroxy-1,1-diphosphonic acid
Perborate	:	Sodium perborate monohydrate
Percarbonate	:	Sodium percarbonate of the nominal formula $2\text{Na}_2\text{CO}_3 \cdot 3\text{H}_2\text{O}_2$
Carbonate	:	Anhydrous sodium carbonate
Termamyl	:	α -amylase available from Novo Nordisk A/S
Savinase	:	protease available from Novo Nordisk A/S
FN3	:	protease available from Genencor
SLF18	:	low foaming surfactant available from Olin Corporation
ACNI	:	alkyl capped non-ionic surfactant of formula $\text{C}_{9/11} \text{H}_{19/23} \text{EO}_8$ -cyclohexyl acetal
C_{14}AO	:	tetradecyl dimethyl amine oxide
C_{16}AO	:	hexadecyl dimethyl amine oxide
Duramyl	:	α -amylase available from Novo Nordisk A/S
DPM	:	dipropylene glycol methyl ether
DPG	:	dipropylene glycol
Methocel	:	cellulosic thickener available from Dow Chemical

5

In the following examples all levels are quoted as parts by weight.

Examples 1 to 4

- 10 The compositions of examples 1 to 4 are introduced in a two compartment layered PVA pouch. The dual compartment pouch is made from a Monosol M8630 film as supplied by Chris-Craft Industrial Products. 17.2 g of the particulate composition and 4 g of the anhydrous composition are placed in the two different compartments of the pouch. The deformability of the exemplified pouches, measured using an Instron material tester
- 15 (following the method described hereinabove) is 23%. The pouch is introduced in the 25

- 5 ml dispenser compartment of a Bosch Siemens 6032 dishwashing machine, the dispenser is closed and the washing machine operated in its normal 55°C program.

Example	1	2	3	4
<u>Particulate composition</u>				
C ₁₄ AO	4.6	4.6		
C ₁₆ AO			4.6	4.6
ACNI	4.6		4.6	
SLF18		4.6		4.6
STPP (anhydrous)	27.5	27.5	27.5	27.5
STPP (hydrated)	27.5	27.5	27.5	27.5
HEDP	1.0	1.0	1.0	1.0
Savinase	1.3	0.7	0.7	1.7
Termamyl	1.7	0.7	0.7	1.3
FN3	1.6			1.6
Perborate		14.2		14.2
Percarbonate	14.2		14.2	
Carbonate	9.3	9.7	9.0	9.3
Silicate	6.4	6.0	6.7	6.4
Perfume	0.3	0.5	0.5	0.3
<u>Anhydrous composition</u>				
DPG	99.5	95.0	95.0	99.5
FN3 Liquid		2.60	2.4	
Duramyl Liquid		2.0	2.4	
Dye	0.5	0.4	0.2	0.5

Examples 5 to 8

- 10 42 g of the compositions of examples 5 to 8 are introduced in a single compartment PVA pouch of 36 mm volume. The pouch is made from a Monosol M8630 film as supplied by Chris-Craft Industrial Products. The exemplified compositions are in the form of a paste having a density of 1300 kg/m³. The deformability of the exemplified pouches, measured

- 5 using and Instron material tester (following the method described hereinabove) is 30%. The pouch is introduced in the 42 ml single compartment dispenser of a Whirlpool dishwashing machine, the dispenser is closed and the washing machine operated in its normal 65°C cycle.

10

Example	5	6	7	8
C ₁₄ AO	0.5	5.6		
C ₁₆ AO			3.6	0.5
ACNI	4.6		4.6	
SLF18		5.6		4.6
STPP (anhydrous)	34	33	36	34
DPM	45.2		45.6	46.1
DPG		45.5		
Savinase		1.7	1.6	1.6
Termamyl	1.6	1.6		
FN3	2.0		1.6	1.0
Carbonate	10	10	10	10
Methocel	0.5	0.5	0.5	0.4
Perfume	0.5	0.5	0.5	0.5

Examples 9 to 12

- 50 g of the compositions of examples 9 to 12 are introduced in a single compartment PVA pouch of 36 mm volume. The pouch is made from a Monosol M8630 film as supplied by Chris-Craft Industrial Products. The exemplified compositions are in the form of a paste having a density of 1300 kg/m³. The deformability of the exemplified pouches, measured using and Instron material tester (following the method described hereinabove) is 30%. The pouch is introduced in the 42 ml single compartment dispenser of a Whirlpool dishwashing machine, the dispenser is closed and the washing machine operated in its normal 65°C cycle.

5

Example	9	10	11	12
C ₁₄ AO	1.2			
C ₁₆ AO		1.2	1.1	
ACNI	1.1	1.1		
SLF18			1.1	
LF404				2.3
STPP (anhydrous)	34	34	34	34
Silicate	8	8	8	8
HEDP	0.7	0.7	0.7	0.7
Percarbonate	10	10	10	10
DPM	30		30	
DPG		30		30
Savinase	0.5	0.5	0.5	0.5
Termamyl	0.5			0.5
FN3		0.5	0.5	
Carbonate	13	13	13	13
Methocel	0.5	0.5	0.5	0.4
Perfume	0.5	0.5	0.5	0.5